



Empowering families.  
Fighting Duchenne.

JETT FOUNDATION'S

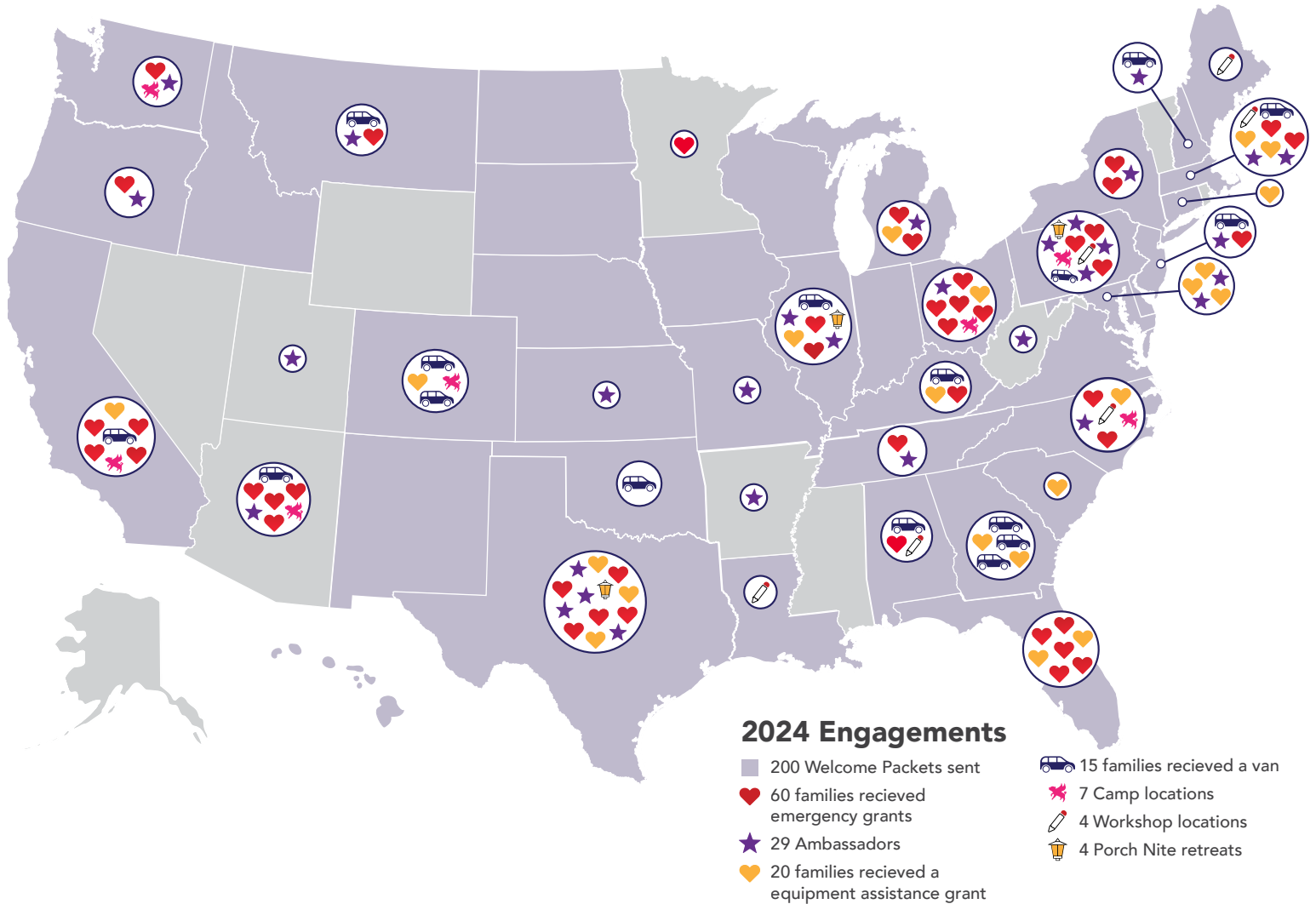
# IMPACT Report

FY24





Jett Foundation empowers people and families impacted by Duchenne muscular dystrophy through the development of transformative programming, educational opportunities, and ongoing support for every stage of a Duchenne journey.



## 2024 HIGHLIGHTS AT A GLANCE

**Served more than 4,000 individuals** (35% more than in 2023).

**261 campers participated** in Camp Promise programs, including 162 young adults who are 18+.

**877 individuals/families** registered for our Family Workshops and Community Webinars, with an additional 1,500 views on YouTube.

**200+ applications** were received for our Jett Giving Fund Accessible Vehicle Fund, Emergency Fund, and Equipment Fund. 120 grants and equipment were awarded.

**250+ individuals** (from 42 different states) requested our Welcome Packets which provide valuable information and resources. Now available in Spanish!

**456 caregivers** registered for our in-person or virtual support groups, including Porch Nite and Carrier Conversations.

**250 participants** in our Team Jett and Signature Events to raise funds and awareness for our programs and the people we serve.

**147,000+ website page views** and 280,000+ reach across social media platforms in 2024.

# Letter from the Executive Director

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Dear friends,

I am filled with immense pride and gratitude as I reflect on the incredible strides Jett Foundation has made in our fight against Duchenne muscular dystrophy and in our unwavering support of the families we served this year. 2024 has been a year of growth, connection, and significant impact, and I am thrilled to share some of the highlights with you.

This year, we continued to build on the momentum of our programs and initiatives, reaching more individuals and families than ever before. Our commitment to providing vital resources, fostering community, and driving progress in Duchenne care remains at the heart of everything we do.

This year also marked a significant milestone as we were able to host our **first in-person World Duchenne Awareness Day event in four years!** It was a truly special occasion, bringing together 150+ members of the community—including families, industry partners, and fellow Duchenne advocacy organizations—for a day of connection, education, and inspiration. Recognizing the importance of inclusivity, we also maintained a robust virtual component, ensuring that individuals from across the globe could participate and feel part of this important day.

Expanding our reach and impact even further, we were excited to launch **two new Camp Promise locations this past summer: Camp Promise Southeast and Camp Promise Southern California.** These additions to our camp lineup provided even more young people and adults living with Duchenne and other neuromuscular disorders with the opportunity to experience the joy, friendship, and independence that Camp Promise offers.

As we look ahead to the coming year, we are filled with excitement and renewed determination. We are committed to expanding our programs, strengthening our partnerships, and continuing to provide unwavering support to families navigating the challenges of Duchenne. We have big plans in store, all aimed at making a tangible difference in the lives of those affected by Duchenne.

Thank you for being an integral part of the Jett Foundation family. Your support, dedication, and belief in our mission make our work possible. **Together, we are truly making a difference.**

With heartfelt gratitude,

**Eric Snyder**  
Executive Director  
Jett Foundation

# Jett Giving Fund Empowers Families with Duchenne Through Essential Support

Launched in 2017, the Jett Giving Fund was established to address the critical needs of families impacted by Duchenne muscular dystrophy, particularly the significant financial burden associated with essential equipment like accessible vehicles. Recognizing the profound impact of limited mobility on daily life, the Jett Giving Fund has expanded its support to include emergency assistance and equipment grants, empowering families to navigate the challenges of living with Duchenne.

The Jett Giving Fund currently offers three main avenues of support:



## 1) The Accessible Vehicle Fund

Assists families in acquiring accessible vehicles, enabling them to maintain independence, access medical care, and participate in community activities.

- **15 families** accepted into the program
- **42 applications** received in 2024
- **2 accepted families** had 2 children with Duchenne, 1 family had 3 children with Duchenne
- **Families** accepted into the program came from 10 different states

## A FAMILY TRANSFORMED: The Cruz Family Story

The impact of the Jett Giving Fund is evident in the inspiring story of the Cruz family from Gainesville, Georgia. Three brothers, Victor (23), Christian (18), and Alan (16), all live with Duchenne muscular dystrophy. Due to the limitations of their existing vehicle, the family faced significant challenges in transporting their sons, often requiring them to manually lift wheelchairs and navigate unreliable public transportation.

*“Traveling with the boys and lifting them each time became unsafe and nearly impossible,”* shares their father, Martin.

Through the Jett Giving Fund’s Accessible Vehicle Fund, the Cruz family received crucial financial assistance to purchase a fully accessible van, significantly improving their mobility and quality of life. With their new vehicle, the family can now travel together safely and comfortably, attend medical appointments, and explore the world with renewed freedom.

*“This accessible van is life-changing for our family,”* says Martin. *“We will now have unencumbered travel, and this will significantly reduce the physical tax on me and my wife’s bodies.”*

The Cruz family’s journey exemplifies the profound impact of the Jett Giving Fund in empowering families affected by Duchenne to navigate the challenges of daily life with greater independence and dignity.





Similarly, the Jett Giving Fund brought much-needed relief to the Dinh family of Dracut, Massachusetts (pictured on cover). Kristina and Si Dinh faced the difficult reality of their sons, Steven (15) and Simon (11), being diagnosed with Duchenne muscular dystrophy. With their sons relying on power wheelchairs and unable to move independently, the simple act of leaving their home became a daunting task. Lifting and carrying Steven and Simon to their vehicle posed a significant safety risk, especially with their young son, Dennis, also needing care.

*"We couldn't safely get the boys to appointments, school, tutoring, and daily activities," shares Kristina and Si. "The boys deserve to see new places and get the care they need."*

By connecting with Jett Foundation and participating in the Accessible Vehicle Fund, the Dinh family found a path to greater freedom. After successfully fundraising half the cost of their vehicle, Jett Foundation, with the support of sponsors, covered the remaining expenses. The Dinh family celebrated the arrival of their new accessible van surrounded by love and support. The day was filled with joy, happy photos, and heartfelt gratitude, marking the end of years of transportation struggles.

*"In the midst of our struggles, we found kindness," Kristina says. "When we were accepted into the Accessible Vehicle Fund, we weren't just offered support; we were offered freedom. At the time, a safe and accessible van was something we could only dream of. Now, our boys can safely go to school, the park, and appointments without struggle."*

### Community Collaborations

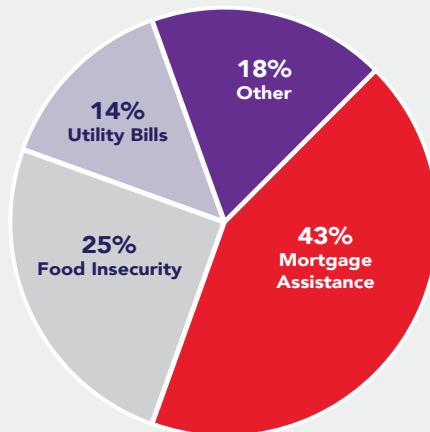
By joining forces with other Duchenne organizations, we were able to provide three more accessible vehicles, and we're incredibly proud of this community effort.



### 2) Emergency Fund

Provides critical financial assistance to families facing unforeseen challenges, such as unexpected medical expenses or home modifications.

- In 2024, we granted **over 80 grants** to families from **21 different states** in the amount **over \$100,000** in the following areas:



*Other: these were for car repairs, childcare, and hurricane/weather-related emergencies*



### 3) Equipment Fund

Supports the acquisition of essential equipment, such as specialized wheelchairs, communication devices, and adaptive technology, to enhance quality of life and independence.

- In the first year of this program, Jett Foundation gifted **15 grants** to families in the amount of **\$30,000**.
  - Types of assistance include: Hoyer lifts, travel chairs, respiratory equipment, ramps, and assistive technology





## JETT FOUNDATION'S CAMP PROMISE: A Summer of Joy and Connection for Individuals with Neuromuscular Disorders

**Jett Foundation's Camp Promise**, a summer camp program for individuals with neuromuscular disorders, successfully completed its 2024 tour with seven exciting locations across the United States, in addition to a virtual weekend. This year, the program expanded its reach with the addition of two new stops: Southern California and South Carolina, providing more opportunities for young people and adults to experience the magic of Camp Promise.

Camp Promise provides a valuable opportunity for individuals with Duchenne and other neuromuscular disorders to connect with peers, build confidence, and create lasting memories. The program fosters a supportive community and empowers campers to explore their abilities and live life to the fullest.

Campers participate in traditional summer camp activities and build confidence through unique programming that encourages them to step out of their comfort zone and engage with others who live with neuromuscular diseases. Camp Promise offers a wide range of activities, including accessible ropes courses, ziplining, archery, swimming, arts and crafts, games, music, and more. Campers engage in themed activities, build friendships, and participate in community-building events, creating lasting memories.





**Camp Promise is all about firsts!**  
**Here are a few highlights**

- Attending Camp Promise
- Catching a Bass fish
- Doing archery
- On a wheelchair accessible porch swing
- On an accessible pontoon boat
- Playing Bingo bingo
- Playing trivia with my friends
- Playing wheelchair soccer
- Racing a boat I made
- Riding an accessible bike!!
- Riding an accessible golf cart
- Singing karaoke
- Smashing an egg on someone's head
- Trying a chicken quesadilla
- Trying Italian dressing
- Trying whole grain cheerios
- Using a hooyer lift at camp
- Winning Bingo
- Ziplining



*"I really enjoyed kayaking. It was definitely challenging but I felt very safe and it was a very rewarding experience."*



**Impactful Reach in 2024:**

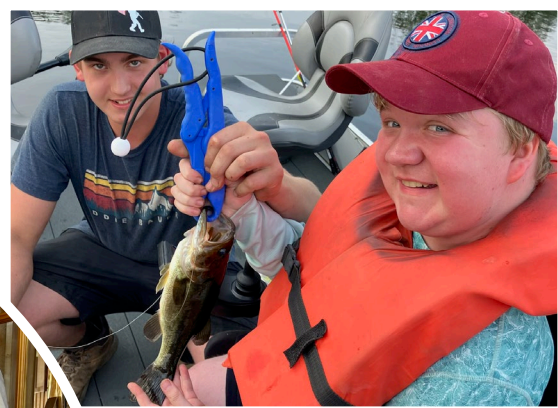
- **Camp Promise served a total of 261 campers** (an increase of over 100 from 2023) at eight sessions in the United States.
  - › **148** in-person Camp Promise participants supported by 200 volunteers
  - › **113** Camp Promise Connect virtual participants
  - › **500+** parents, siblings, and caregivers provided an opportunity for respite
- **62%** of camp participants served are ineligible to attend any other camp due to age restrictions or the specific needs of their condition

**Beyond Camp**

Camp Promise extends beyond the summer months, fostering a sense of community and support year-round. Campers develop lasting friendships and connections, providing a valuable support system throughout the year with virtual programs and events.

**Jett Foundation's Camp Promise continues to be a beacon of hope and joy for individuals with neuromuscular disorders, empowering them to live full and fulfilling lives.**







**Jett Foundation is committed to empowering families and individuals affected by Duchenne muscular dystrophy** through comprehensive community education programs. Jett Foundation provided ongoing support through two key programs in 2024: Meet and Greet and Welcome Packet programs.

## **MEET AND GREET PROGRAM: Building Connections Across Communities**

**Jett's Meet and Greet program** utilizes one-hour Zoom meetings to connect with clinics, schools, DSS agencies, and others, particularly those in rural and southern states where diagnosis delays are prevalent. The goal is to share Jett Foundation's resources and educational materials with both clinicians and patients. In 2024, Jett Foundation met with 18 clinics, schools, DSS agencies, and others that collectively **serve over 500 patients annually**. Participating institutions included, but were not limited to:

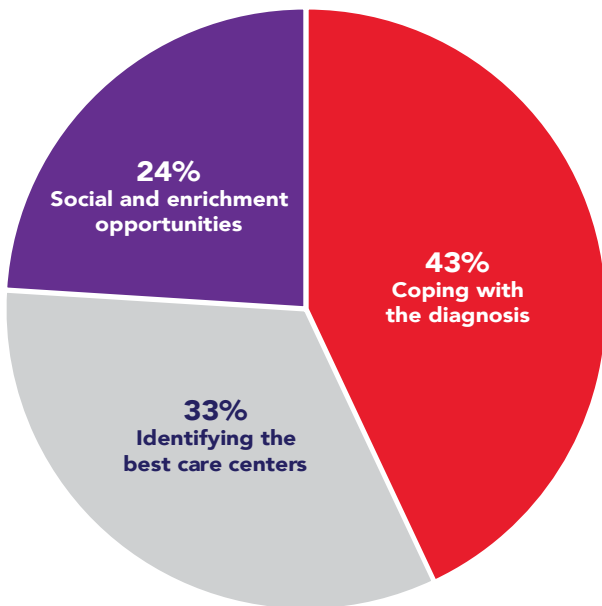
- University of Alabama, Birmingham, AL
- Jones County Department of Social Services, Trenton, NC
- Saint Anthony's Group, MDA Care Center, Saint Petersburg, FL
- John's Hopkins Children's Hospital, Saint Petersburg, FL
- Nonotuck Resource Associates, Inc. Plymouth, MA
- Charlotte-Mecklenburg Schools, Charlotte, NC
- Nation Wide Children's Hospital, Columbus, OH
- Akron Children's Hospital, Akron, OH
- Hackensack Meridien Health, Hackensack, NJ
- Norton Health, Louisville, KY



## WELCOME PACKET PROGRAM: A First Point of Contact

Jett Foundation's **Welcome Packets** serve as a crucial first touchpoint for many families navigating a Duchenne diagnosis. It has also become a trusted resource for caregivers, clinicians, industry partners, and other key stakeholders. The packet contains valuable information on treatment, clinical trials, support letters from the community, and details about Jett Foundation's programs.

In 2024, **over 250 Welcome Packets** were requested by individuals in **42 states**. Families requesting packets identified the following as areas of support they were interested in the most:



These initiatives demonstrate Jett Foundation's commitment to providing accessible education and support to the Duchenne community, empowering families with the knowledge and resources they need to navigate their journey.

## Family Workshops

Jett Foundation's commitment to supporting families affected by Duchenne muscular dystrophy extends into vital educational programming. Jett's Family Workshop series brings together leading clinicians, researchers, and families to share knowledge, discuss cutting-edge therapies and clinical trials, and connect families with valuable local resources. This national education program provides a crucial link between families and the latest advancements in Duchenne care.

These workshops offer more than just information; they foster a sense of community and shared understanding, empowering families with the knowledge and resources they need to navigate the complexities of Duchenne. By connecting families with leading experts and local resources, Jett Foundation strengthens the support network for those affected by this challenging condition.

These intimate meetings offer a unique opportunity for families to learn from experts in a supportive environment. In 2024, Jett Foundation hosted four regional workshops, impacting over **150 registered participants**.



*// It was so inspiring to hear directly from families and clinicians – there was a real sense of community and safety in the room”*

### The 2024 workshop locations included:

- **LCMC Health**  
New Orleans, LA (April 6, 2024)
- **Levine's Children's Hospital**  
Charlotte, NC (May 4, 2024)
- **Maine Health**  
Portland, ME (September 28, 2024)
- **Penn State Health**  
Harrisburg, PA (November 16, 2024)

# Webinars

Jett Foundation’s commitment to supporting the Duchenne muscular dystrophy community extends far beyond in-person events. The organization’s Community Webinar Series provides an accessible and convenient way to bring vital educational resources directly into the homes of families and individuals affected by Duchenne, no matter where they live.

These webinars offer a unique opportunity for participants to connect with and learn from leading experts in a variety of crucial areas. From practical tips for daily care to updates on cutting-edge technology and treatments, the series covers a wide range of topics relevant to the Duchenne community. Participants can not only hear from experts but also engage in live Q&A sessions, ensuring their specific questions and concerns are addressed.

The impact and value of the Community Webinar Series are clearly reflected in participant feedback:

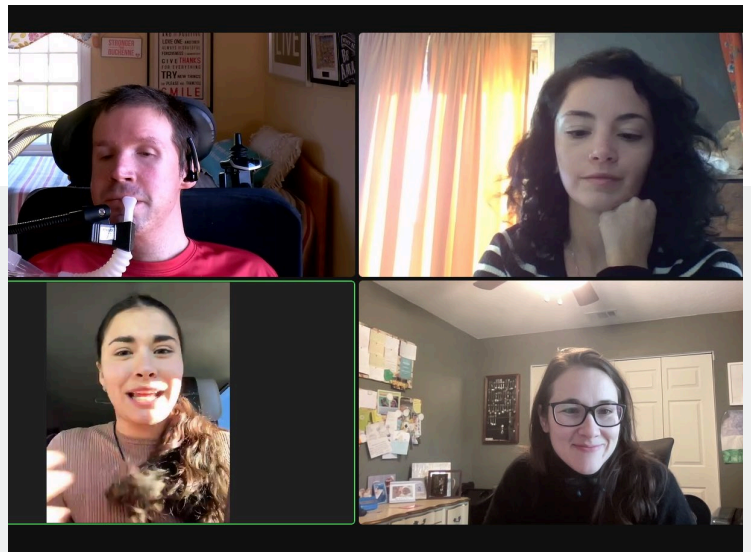
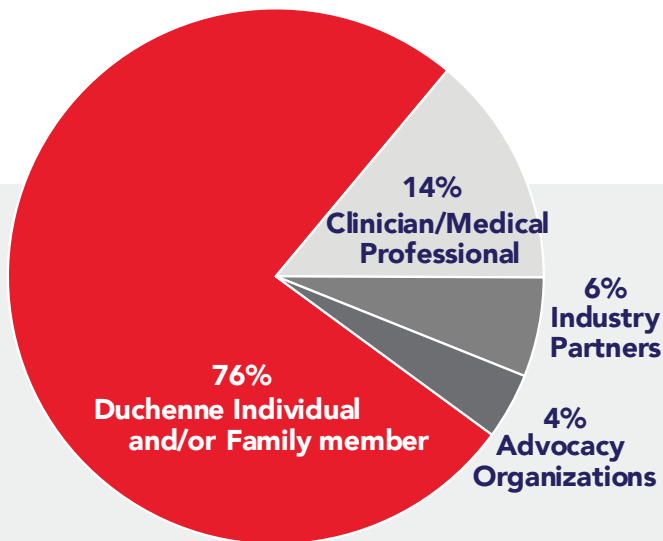
- **96% of participants** indicated they were “very likely” to recommend the webinar to someone impacted by Duchenne or seeking information on the topic.
- **100% of participants** expressed they were “very likely” to attend another Community Webinar.

These overwhelmingly positive responses underscore the effectiveness of the series in providing valuable and accessible education to the Duchenne community. Jett Foundation’s Community Webinar Series serves as a vital resource, empowering families and individuals with the knowledge they need to navigate the complexities of Duchenne and live their fullest lives. By offering these educational opportunities online, Jett Foundation ensures that geographical limitations do not become barriers to accessing critical information and support.

In 2024, Jett Foundation hosted **22 webinars**, demonstrating the program’s significant reach and impact. Over **400 individuals registered** for these webinars, representing an impressive **42 states** within the United States and **11 countries** across the globe: Poland, India, Romania, Switzerland, Sweden, Australia, Canada, Korea, Malaysia, England, and Egypt. This diverse participation highlights the global need for Duchenne education and the accessibility of Jett Foundation’s resources.

The impact of these webinars extended beyond live attendance. An additional **1,500 views of recorded webinars** were logged through Jett Foundation’s website and YouTube page, demonstrating the enduring value and accessibility of this educational content. Facebook Live streams consistently increased webinar viewership by an average of **300 views per event**. This on-demand access ensures that individuals can benefit from the expertise shared in the webinars at their convenience.

2024 WEBINAR REGISTRANTS





## COMMUNITY AMBASSADORS: A Vital Force

Our **Community Ambassadors** are invaluable members of the **Jett Foundation family**. Often the first point of contact for newly diagnosed families, these dedicated volunteers play a crucial role in welcoming families into the community through regional meetups and virtual panel discussions. In 2024, **31 volunteer Ambassadors** from **23 states** generously shared their time and personal experiences with over **750 members** of our community.

*Key contributions from our Ambassadors this year include:*

- **Industry Advocacy:** Connecting with industry partners to share the valuable perspectives of patients and families.
- **Community Feedback:** Regularly meeting with Jett Foundation staff to provide valuable insights and feedback gathered within the community.
- **Community Engagement:** Actively sharing Jett Foundation announcements, industry news, events, and webinars within their networks.

*“Jett Foundation is a family-friendly networking resource. It provides social connections/educational opportunities for caregivers/families. Jett reduces stress with financial support to DMD families. They focus on day-to-day functions and let families know they have a Jett community available to them in many ways.”*

## Supporting Families Through Community

**Jett Foundation’s support groups** were born from the recognition of the critical need for connection and support within the Duchenne community. Recognizing the isolation often felt by parents and caregivers, these groups provide a safe and understanding space for families to connect and find solace in shared experiences.

Our support groups have expanded to include valuable resources like Carrier Conversations, specifically designed to support women who are carriers of the Duchenne mutation. These conversations provide a platform for carriers to discuss their fears, hopes, and concerns, while learning how to care for their own health and navigate the emotional and practical challenges associated with being a carrier.



- We hosted **10 virtual Carrier Connections meetings**, which drew a total of **142 registrants**.
- Hosted **14 virtual Porch Nights** which drew a total of **273 registrants**.
- **3 In-person porch nites** (TX, PA, IL), **29 attended**, and 41 registered.



**30 Ambassadors** from 21 states have a combined network of 750 families impacted by Duchenne



**Jessica with her son, Javon**



**Jessica Campbell, MSW (left)** at a Jett Foundation Family Workshop

## STRENGTHENING FAMILY RESOURCES: Jett Foundation Hires Social Worker

Jett Foundation welcomed its first full-time social worker this year, Jessica Campbell, to better serve families impacted by Duchenne muscular dystrophy. Jessica, a mother of two with a son living with Duchenne, brings both professional expertise and personal understanding to her new role. She dedicates her time to helping families navigate the complexities of Duchenne by connecting them with crucial resources, providing emotional support, and being readily available on a full-time basis. This significant step underscores Jett Foundation's commitment to holistic family support, ensuring families have the guidance and assistance they need throughout their journey.

# Jett Foundation Celebrates Awareness with Community Events

Jett Foundation marked two significant awareness days in 2024 with impactful events that brought together the Duchenne community and beyond.





## Rare Disease Day

On February 29th, Jett Foundation celebrated its seventh annual “Thriving with Duchenne” Rare Disease Day event. This year’s event featured a panel discussion moderated by Kyle Bryant and Sean Baumstark, hosts of the “Two Disabled Dudes” podcast, who shared their experiences living with Friedreich’s ataxia.

The Duchenne community was represented by panelists, including Colin Werth, Kris Napper, Race Martinez, Amanda Becker, and Chris Schlechty, who shared their personal journeys of overcoming challenges and achieving

their goals. From navigating college life to achieving driving independence, the panelists offered valuable insights and encouragement to the audience.

Leading up to the event, Jett Foundation invited individuals and families from the rare disease community to submit videos showcasing how they are “living to the fullest.” These inspiring videos were shared on Jett Foundation’s Facebook page, further amplifying the voices of individuals living with rare diseases.

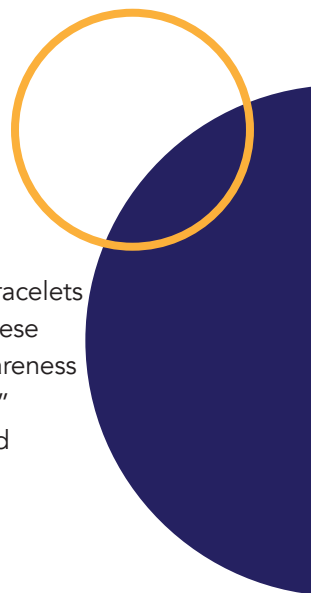
## World Duchenne Awareness Day

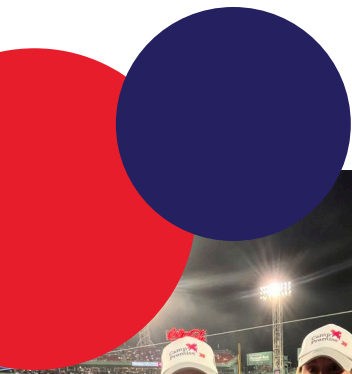
On September 7th, Jett Foundation hosted a series of events to commemorate World Duchenne Awareness Day. The highlight was an in-person luncheon and family event attended by over 150 individuals, including families affected by Duchenne, medical professionals, and Jett Foundation supporters. The event featured engaging activities, a keynote address from Sam Safford, and opportunities for community members to connect. Sam’s remarks emphasized the importance of creating an inclusive society where accommodations are not exceptions but the norm.

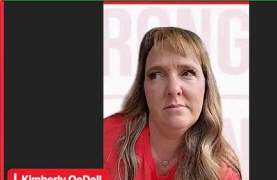
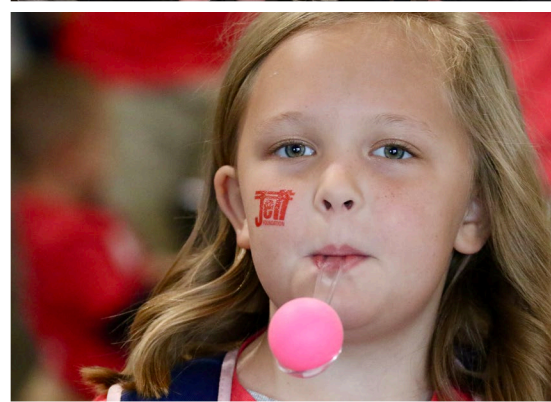
In the evening, Jett Foundation hosted a virtual panel discussion titled “Raise Your Voice for Duchenne,” featuring experts in disability rights and advocacy.

The panel provided valuable insights into the importance of community advocacy and the power of collective action within the Duchenne community.

To further amplify awareness, Jett Foundation distributed 255 awareness boxes containing various items such as window clings, pinwheels, and light-up bracelets to families throughout the community. These boxes empowered families to spread awareness and show their “Stronger than Duchenne” pride within their own neighborhoods and communities.









## Team Jett & Signature Events Drive Impact through Athletic Endeavors

In 2024, Jett Foundation witnessed an outpouring of support from its dedicated community of athletes and event participants. Over 250 individuals joined the Team Jett movement, embracing a variety of athletic challenges to raise crucial funds for families impacted by Duchenne muscular dystrophy.

From exhilarating triathlons like the Cohasset Triathlon and Hopkinton Title 9 Triathlon (Gals for Cal) events to the iconic Falmouth Road Race, Team Jett members demonstrated incredible dedication and perseverance. Their efforts, alongside the inspiring participation of numerous individuals in Jett's annual National Challenge, showcased the power of collective action in making a tangible difference.

The Jett Golf Classic, a signature fundraising event, proved to be a resounding success. With the generous support of **156 participants** and dedicated sponsors, the event **raised \$145,000**. The impact of this fundraising was poignantly highlighted at the post-golf reception, where a grateful family recipient from the Jett Giving Fund's Accessible Vehicle fund shared their inspiring story and showcased their new accessible van.

These remarkable achievements underscore the unwavering commitment of the Jett Foundation community. Through their dedication and generosity, participants are not only supporting vital programs but also empowering families and individuals affected by Duchenne to live more fulfilling lives with greater autonomy and control.





# Thank you to our donors

Our donors make it possible for Jett Foundation to empower families and fight Duchenne. Thank you to the people and organizations who contributed \$1,000 or greater during FY24 through financial gifts, in-kind donations, or peer-to-peer fundraising.



American Plumbing and Heating  
Anderson Fuel  
Arizona Community Foundation  
James Armitstead  
Auburn Construction  
Austin's Hope  
Avidity Biosciences  
Beth Baird  
Ball Family  
Stephanie Bandzak  
Barbara Bresnan Memorial Foundation  
Barkowskie Family  
Barry Carroll Foundation for Lodge 25 IN  
Bay State Federal Savings Charitable Foundation  
Bergen Count Sheriff's Department  
Beth Israel Deaconess Hospital  
Biomarin  
Bob's Discount Furniture Charitable Foundation, Inc.  
Nicole Borden  
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Scott Clifford  
Coastal Bend Community Foundation  
Amanda Coleman  
Stephanie Colorado  
Lora Cooley  
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Crossfit Mob  
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Deen Day Sanders  
Amanda Dehart  
Sheila and Tony Dicioccio  
Joanne Donovan  
Dr. Christiane Rougee Memorial Fund  
Dr. Squatch  
Kevin Ducie  
Dyne Therapeutics  
Edgewise Therapeutics  
Emilson Family Foundation  
Entrada Therapeutics  
Family Nail Inc  
Jim Ferguson  
Emily Fields  
Finn's Friends  
Firefighters of PBC  
Firefighters on a Mission  
Focus

## CHAMPIONING OUR MISSION: Cure Dale's Duchenne

Cure Dale's Duchenne demonstrates a profound commitment to supporting families facing the challenges of Duchenne muscular dystrophy. Having personally navigated Duchenne with Dale, their son, they offer meaningful assistance rooted in deep understanding and compassion. In 2017, Cure Dale's Duchenne became a vital partner of the Jett Giving Fund, providing crucial support to our Accessible Vehicle Fund. With a spirit of humility and a genuine desire to positively impact the lives of families impacted by Duchenne, their annual

support has been instrumental in empowering families with the freedom and independence that accessible vehicles provide.

As Lelia Ginder, co-founder of Cure Dale's Duchenne, explains, "We were deeply moved by the number of families who struggled to provide the necessary support for their children. Navigating the challenges of Duchenne is incredibly difficult, and the financial burden of essential equipment like accessible vehicles can be overwhelming."

Rick Ginder, co-founder of Cure Dale's Duchenne, echoes this sentiment: 'Jett Foundation has a well-established reputation for transparency and genuine support within our community. We trust in their mission and believe that by partnering with Jett, we can maximize our impact and ensure that families have access to the critical resources they need.' By choosing to support Jett's existing programs, Cure Dale's Duchenne demonstrates a commitment to efficiency and a deep understanding of the urgent needs within our community."

Fraternal Order Of Police Lodge  
 FRC HOLDCO, LLC  
 John Furth  
 Kenneth Gaaney  
 Ganot Family  
 Laurie and John Garbarino  
 The Ginder Family  
 Glance Networks  
 Global DMC Partners  
 Global Vision Advisors  
 Alexis Grace  
 Steven and Debbie Greenwall  
 Hale Family Foundation  
 Martha Hanson  
 Scott Hastings  
 Jamie Hawes and Ellen Hanson  
 Help Hope Live  
 Heritage Print Solutions  
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 Law Offices of Lipsey & Clifford  
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 Craig Lipman  
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 Scott MacDonald  
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 Mahoney Law Group, P.C.  
 Peter Mair  
 Mammoth LLC  
 Craig Martin

Mo McDonnell  
 McSherry Family  
 Michael's Cause Inc.  
 MT Olive UCC  
 Mt. Pleasant Baptist Church  
 My DMD Hero Inc.  
 Dzu Nguyen  
 Le Nguyen  
 Long Nguyen  
 Peter Nguyen  
 Nicholoff Family  
 NS Pharma  
 Numotion Foundation  
 Ocean Properties  
 Oceanside Restoration  
 Jeananne Oliphant  
 Optio Biopharma Solutions  
 Scott Oran  
 Robert E & Marie Orr Smith  
 Foundation  
 Orsini Pharmaceutical Services  
 Tom Osuch  
 Nicolas Pakler  
 Pathway Community Church  
 Jennifer Patten  
 PBC Benevolent Fund  
 PepGen Inc  
 Percheron Therapeutics  
 Peters Family Foundation  
 Jim and Lynda Petropulos  
 Pfizer Inc.  
 Pledgeling Foundation  
 Erin Presutti  
 PTC Therapeutics  
 Purchasing Partners, Inc  
 Quincy Mutual Fire Insurance  
 Quitzau Family  
 Regenxbio  
 Diane Repress  
 Retail Storefront Group  
 Leo Robinson  
 Rodman Ride for Kids  
 Romito Foundation  
 William Rose  
 Gilbert Rosenberg  
 Jacob Rosenberg  
 Michael Ryan  
 Ryan's Quest  
 Sarepta Therapeutics, Inc.

Satellos  
 Scalora Consulting Group  
 Shea Concrete Products, Inc.  
 Chris and Dawn Shee  
 John Skubina  
 Smith Sullivan & Brown PC  
 Yancey Snavelly  
 Eric and Melanie Snyder  
 Solid BioSciences  
 South Shore Bank  
 Christine Stalling  
 Laurie Stalling  
 Gregory Stratis  
 Cindy Sullivan  
 The Suneel Research  
 Foundation  
 Robert Thal  
 The Assistance Fund  
 The Everyday Good Inc  
 The Family Office at Synovus  
 The Flatley Foundation

The HCA Foundation  
 The Yale Charitable  
 Foundation  
 Tito's Handmade Vodka  
 TLC Pediatrics  
 Transamerica  
 Lynn Treadaway  
 Matt Trudeau  
 John Turco  
 Ultragenyx Pharmaceutical Inc.  
 Vertex Pharmaceuticals  
 Wave Life Sciences  
 Mrs. Mary-Lou Weisman  
 Wellington Management  
 WestRock  
 William Wheaton  
 Wheels for Wheels  
 Maxline & Wilson Family  
 Woodland Partners  
 Suzanne York  
 Christine Zinman

## FY24 Financials

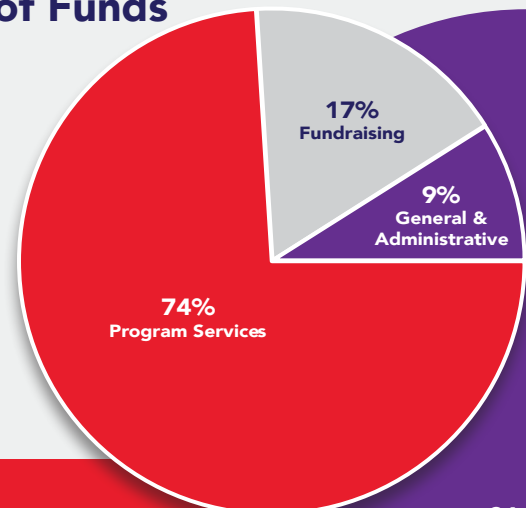
### OPERATING REVENUE

Foundation & Corporations	\$ 2,624,276
Individuals Contributions:	\$ 1,443,378
Fundraising Events	\$ 215,954
<b>Total Support</b>	<b>\$ 4,283,608</b>

### OPERATING EXPENSES

Program Services	\$ 2,504,374
Fundraising	\$ 593,835
General & Administrative	\$ 324,470
<b>Total Expenses</b>	<b>\$ 3,422,679</b>

## Use of Funds



## STAFF

Eric Snyder, Executive Director

## FINANCE & HR

Allison Caputo

## PROGRAMS

Niki Benfield

Jessica Campbell

Meghan Houston

Julie LeClair

Trina Stelly

Alexa Tinsley

## DEVELOPMENT

Maura Carroll

Emily Hubert

## MARKETING & COMMUNICATIONS

Zac Bentley

Gabriella Costa

## BOARD OF DIRECTORS

### Executive Board:

**Christine McSherry**, Jett Foundation Founder

**Ellen Hanson, Vice-Chair**, Boston Children's Hospital

**Alfred Jackson, Chair**, Ketchum, Inc.

**Michael Mullin, Treasurer**, Claro Advisors, LLC

### Directors:

**Diana Johnson**, Elite Clinical Network

**Adrian Kielhorn**, Alexion Pharmaceuticals

**Melissa Paige**, National Association of Medication  
Access & Patient Advocacy

**Colleen Pike**, Lhasa OMS

**Leo Robinson**, Wells Fargo Advisors

**Chris Schlechty**, Microsoft

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